Claims:

- 1. (Original) A method of detaching a nucleic acid molecule from a solid support to which it is attached, wherein an unconventional nucleotide is incorporated at a predetermined site in said nucleic acid molecule, said method comprising selectively cleaving said nucleic acid molecule at the site of said unconventional nucleotide, wherein said selective cleavage is accomplished enzymically.
- 2. (Previously amended) A method of reversibly immobilising a nucleic acid molecule, said method comprising:
- (a) incorporating an unconventional nucleotide into said nucleic acid molecule at a pre-determined site;
- (b) binding said nucleic acid molecule to a solid support; steps (a) and (b) being carried out in either order or simultaneously; and subsequently
- (c) selectively cleaving said nucleic acid molecule at the site of said unconventional nucleotide, wherein said selective cleavage is accomplished enzymically.
- 3. (Original) A method as claimed in claim 1 or claim 2 wherein said nucleic acid molecule is a chimeric molecule comprising a nucleic acid component and another non-nucleic acid component.
- 4. (Previously amended) A method as claimed in claim 1 or 2, wherein the unconventional nucleotide is uracil, hypoxanthine, a ribonucleotide, –7 methylguanine, 8-oxoguanine, deoxyuridine, deoxyinosine, deoxy 5,6-dihydroxythimine, 5'6'-dihydroxythine, deoxy 3'-methyladenosine or 3'-methyladenosine.
- 5. (Previously amended) A method as claimed in claim 1 or 2, wherein said selective cleavage is achieved using a DNA glycosylase enzyme.
- 6. (Previously amended) A method as claimed in claim 1 or 2, wherein said nucleic acid molecule comprises DNA, said unconventional nucleotide is uracil (U), and selective cleavage is achieved using a uracil DNA glycosylase enzyme (UDG).

- 7. (Previously amended) A method as claimed in claim 1 or 2, wherein said unconventional nucleotide is incorporated into said nucleic acid molecule as part of a linker sequence.
- 8. (Original) A method as claimed in claim 7 wherein said linker sequence is a primer.
- 9. (Previously amended) A method as claimed in claim 1 or 2, wherein said nucleic acid molecule is a primer extension product.
- 10. (Previously amended) A method as claimed in claim 1 or 2, wherein said support is a magnetic bead.
- 11. (Previously amended) A method as claimed in claim 7, wherein said linker sequence is provided with means for immobilization to a solid support.
- 12. (Previously amended) A method as claimed in claim 9, wherein said nucleic acid molecule is a cDNA, or a product of an *in vitro* amplification reaction or a sequencing reaction.
- 13. (Previously amended) A method as claimed in claim 7, wherein said nucleic acid molecule comprises a linker sequence coupled to a protein, an enzyme substrate, a receptor ligand, an antigen or hapten, or a fragment thereof, or to an affinity binding group or a reporter group.
- 14. (Currently amended) A method of preparing a construct for binding to, and subsequent cleavage from, a solid support, said method comprising incorporating into said construct a nucleotide linker sequence comprising at a predetermined site an unconventional nucleotide capable of selective cleavage using an a glycosylase enzyme.
- 15. (Currently amended) A chimeric molecule comprising a nucleotide linker sequence comprising at a pre-determined site an unconventional nucleotide capable of selective cleavage using an a glycosylase enzyme, coupled to a functional group.
- 16. (Original) A chimeric molecule as claimed in claim 15, wherein said functional group is an affinity binding group or a reporter group.

- 17. (Previously amended) A method as claimed in claim 14, wherein said linker sequence is immobilized or provided with means for immobilization to a solid support.
- 18. (Previously amended) A chimeric molecule as claimed in claim 16 wherein said affinity binding group is an antibody or a fragment or derivative thereof, or a hapten.
- 19. (Currently amended) A method for separating a target cell from a sample, said method comprising binding said target cell to a solid support by means of a chimeric molecule comprising a nucleotide linker sequence comprising a selectively cleavable unconventional nucleotide at a pre-determined site, coupled to a functional group, preferably as defined in claim 15, wherein said functional group is an affinity binding group which binds specifically to said cell.
- 20. (Currently amended) A method of detaching a nucleic acid molecule from a solid support to which it is attached, wherein an unconventional nucleotide is incorporated a predetermined site in said nucleic acid molecule, said method comprising selectively cleaving said nucleic acid molecule at the site of said unconventional nucleotide, or of reversibly immobilizing a nucleic acid molecule, said method comprising:
- (a) incorporating an unconventional nucleotide into said nucleic acid molecule at a pre-determined site;
- (b) binding said nucleic acid molecule to a solid support; steps (a) and (b) being carried out in either order or simultaneously; and subsequently
- (c) selectively cleaving said nucleic acid molecule at the site of said unconventional nucleotide, preferably as claimed in claim 1 or 2, or a method as claimed in claim 19,

wherein a multiplicity of different nucleic acid molecules or chimeric molecules comprising a nucleotide linker sequence comprising a selectively cleavable unconventional nucleotide at a pre-determined site, coupled to a functional group, are attached or bound to a solid support, each said different molecule incorporating a different unconventional nucleotide.

- 21. (Previously amended) A kit for use in a method as defined in claim 1 or 2, said kit comprising
- (a) means for introducing an unconventional nucleotide into a nucleic acid molecule; and
- (b) means for selective cleavage of said unconventional nucleotide, wherein said means is an enzyme.
- 22. (Currently amended) A poly- or oligonucleotide incorporating an unconventional nucleotide which is selectively cleavable using an a glycosylase enzyme, immobilized on a solid support or carrying means for immobilization.
- 23. (Original) A poly- or oligonucleotide as claimed in claim 22, being poly- or oligo dU.
 - 24. (Original) A poly- or oligonucleotide according to claim 22 being a primer.
- 25. (Previously amended) A poly- or oligonucleotide as claimed in claim 22, wherein said means for immobilization is biotin.
- 26. (Previously amended) A poly- or oligonucleotide as claimed in claim 22, wherein said solid support comprises magnetic beads.
- 27. (Previously amended) A multiplicity of olig- or polynucleotides as defined in claim 22, wherein each different oligo- or polynucleotide incorporates a different unconventional nucleotide.
- 28. (Currently amended) A method as claimed in a chimeric molecule as claimed in claim 15 or 16, wherein said linker sequence is immobilized or provided with means for immobilization to a solid support.
- 29. (New) A method for separating a target cell from a sample, said method comprising binding said target cell to a solid support by means of a chimeric molecule comprising a nucleotide linker sequence comprising a selectively cleavable unconventional nucleotide at a pre-determined site, coupled to a functional group, as defined in claim 15, wherein said functional group is an affinity binding group which binds specifically to said cell.